FORM PTO-1449

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTY, DOCKET NO. VPI/94-04 CIP2 DIV5 SERIAL NO. 09/886,773

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

APPLICANT
Guy W. Bemis, et al.

GROUP 1653

FILING DATE June 21, 2001

U.S. PATENT DOCUMENTS

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EXAMINER INITIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	IF
						APPROPRIATE
200	4,276,298	06/30/81	Jones et al.	424	270	
1 × ×	4,369,183	01/18/83	Jones et al.	424	263	
200	4,499,295	02/12/85	Mueller et al	560	53	
N. DX	4,551,279	11/05/85	Mueller et al.	260	404	
SX A	4,584,397	04/22/86	Mueller et al.	560	75	
S. DL	4,968,607	11/06/90	Dower et al.	435	69.1	
XX OI	5,008,245	04/16/91	Digenis et al.	514	18	
0.2/2	5,055,451	10/08/91	Krantz et al.	514	19	
XX 0.	5,081,228	01/14/92	Dower et al.	530	35.1	
	5,158,936	10/27/92	Krantz, et al.	514	19	
VX O	5,180,812	01/19/93	Dower et al.	530	351	
0125	5,374,623	12/20/94	Zimmerman et al.	514	17	08/20/92
Dit o	5,411,985	05/02/95	Bills et al.	514	460	05/17/93
0,000	5,416,013	05/16/95	Black et al.	435	226	02/18/04
DE O	5,430,128	07/04/95	Chapman et al.	530	330	11/21/94
V X X	5,434,248	07/18/95	Chapman et al.	530	330	06/02/93
W n	5,462,939	10/31/95	Dolle et al.	514	231.5	05/07/93
~ # # P	5,486,623	01/23/96	Zimmerman et al.	549	417	12/08/93
DX O	5,498,616	03/12/96	Mallamo et al.	514	300	11/04/94
~ XX	5,498,695	03/12/96	Daumy et al.	530	331	12/12/94
82 n	5,552,400	09/03/96	Dolle et al.	514	221	06/08/94
A Ald	5,565,430	10/15/96	Dolle et al.	514	19	08/02/94
A CONTRACTOR OF THE PARTY OF TH	5,585,357	12/17/96	Dolle et al.	514	18	01/29/96
No Al	5,585,486	12/17/96	Dolle et al.	544	182	05/12/95
W O A	5,639,745	06/17/97	Dolle et al.	514	183	05/25/95
a de	5,670,494	09/23/97	Dolle et al.	514	86	11/20/95

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David

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FORM PTO-1449 SERIAL NO. U.S. DEPARTMENT OF COMMERCE ATTY. DOCKET NO. PATENT AND TRADEMARK OFFICE VPI/94-04 CIP2 DIV5 09/886,773 **APPLICANT** INFORMATION DISCLOSURE Guy W. Bemis, et al. STATEMENT BY APPLICANT **FILING DATE GROUP** June 21, 2001 1653 EODEIGN DATENT DOCLIMENTS

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1 22	WO 93/25683	12/23/93	PCT	C12N	-15/42		
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W.Z	WO 93/25694	12/23/93	PCT	C12N	15/57		
	WO 94/00154	01/06/94	PCT	A 81K	3 9/89 5		
	WO 94/03480	02/17/94	PCT	G07K	5/02		
WX O 1	WO 95/00160	01/05/95	PCT	A61K-	37/02		
a Wy	WO 95/05192	02/23/95	PCT	-∧61K	38/06		
A O	EP-A-0 275 101	07/20/88	EPO	2871 K	5/02		
7	EP-A-0 410 411	01/30/91	EPO	C07K	-5/04		
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	EP-A-0 479 489	04/08/92	EPO	C07K	-5/93	·	
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	EP-A-0 525 420	02/03/93	EPO	C07D	307/56		
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	OTHER DOCUME		· · · · · · · · · · · · · · · · · · ·			
EXAMINER INITIAL	Principle And Const.	3 17 317355	er Prosert		AL AREA	
Sol	D. Alberg & S. Schreiber, Science, 262, pp. 248-25		sed Design of a	a Cyclophilin-C	Calcineurin Bridgi	ng Ligand",
J.	P. Andrews, "Functional (Sci., 7, pp. 148-151 (1986		Receptor Intera	ections and Dru	ug Design", <u>Tren</u>	ds Pharmacol.
W.	K. Appelt et al., "Design of J. Med. Chem., 34, pp. 19			rative Protein	Crystallographic	Analysis",
22	M. Ator, "Peptide and No Healthtech Institute (Infla					
DX N	M.A. Ator and R.E. Dolle, Curr. Pharm. Design, 1, p			nzyme: Biology	y and the Chemis	stry of Inhibitors"
A WI	E. Baker & J. Drenth, "The Thiol Proteases: Structure and Mechanism", in <u>Biological Macromolecules</u> and <u>Assemblies</u> , 3, pp. 313-368 (F. Jurnak & A. McPherson eds., 1987)					
I O	J. Baldwin et al., "Thienothiopyran-2-sulfonamids: Novel Topically Active Carbonic Anhydrase Inhibitors for the Treatment of Glaucoma", <u>J. Med. Chem.</u> , 32, pp. 2510-2513 (1989)					
W.C.	M. Barinaga, "Death Gives Birth to the Nervous System, But How?", Science, 259, pp. 762-763 (1993)					
	P. Bartlett et al., "CAVEA Molecules", Molecular Re					
DA DY	P. Bender & J. Lee, "Pha pp. 185-193 (1989)	rmacological N	Modulation of In	terleukin-1", A	nnu. Rep. Med. (Chem., 25,
Mr.	R. Black et al., "Activation (1989)	n of Interleukin	-1β by a Co-inc	luced Protease	e", <u>FEBS Lett,</u> 24	7, pp. 386-390
W.	H. Böhm, "The Computer J. Comput. Aided Mol. De			od for the De N	Novo Design of E	nzyme Inhibitors",
AL.	J. Breitner et al., "Inverse Results of a Co-twin Con	Association o	f Anti-inflamma			's Disease: Initial
A.J.	B. Brooks et al., "CHAF Calculations", J. Comput,	RMM: A Prog	ram for Macro	molecular En		on, and Dynamics
W.	A. Brünger, "Extension of Correlation Refinement",	Molecular Re	placement: A N	lew Search Sti	ategy Based on	Patterson
D. D.	A. Brünger & A. Krukowsl Annealing", <u>Acta Crystall</u>	ki, "Slow-Cooli	ing Protocols fo	r Crystallogra	ohic Refinement (by Simulated
	U. Burkert & N. Allinger, " pp. 59-78 (1982)				Geometry" in <u>Mol</u>	lecular Mechanics.
FXAMINER	() 1 D	<u> </u>		באח	F CONSIDERE	6/13/03

FORM PTO-14	449 U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY. DOCKET NO. VPI/94-04 CIP2 DIV5	SERIAL NO. 09/886,773		
IPE	INFORMATION DISCLOSURE	APPLICANT Guy W. Bemis, et al.			
JAN 15 2007	STATEMENT BY APPLICANT	FILING DATE June 21, 2001	GROUP 1653		
P. JAN	OTHER DOCUMENTS (Including Author, Title, I	Date, Pertinent Pages, Etc.)			
EXAMINER INTIAL					
	M. Carson, "Ribbons 2.0", J. App. Crystallogr., 24, pp.	958-961 (1991)			
Il	F. Casano et al., "The Structure and Complete Nucleot Interleukin-1β Convertin Enzyme (ICE)", Genomics, 20	tide Sequence of the Murine Ge	ne Encoding		
DL ,	D. Cerretti et al., "Molecular Cloning of the Interleukin- pp. 97-100 (1992)	*,	<u>∍,</u> 256,		
and the second	K. Chapman, "Synthesis of a Potent, Reversible Inhibit Med. Chem. Lett., 2, pp. 613-618 (1992)	or of Interleukin-1β Converting I	Enzyme", <u>Bioorg.</u>		
Da Ca	N. Cohen, "Drug Design in Three Dimensions", Advance	ces in Drug Research, 14, pp. 4	1-145 (1985)		
	N. Cohen, "Rational Drug Design and Molecular Model		.,		
X î	N. Cohen et al., "Molecular Modeling Software and Met 33, pp. 883-894 (1990)				
Jel Jel	D. Davies & D. Segal, "Protein Crystallization: Micro Techniques Involving Vapor Diffusion", Methods Enzymol, 22, pp. 266-269 (1971)				
De la	K. Dill, "Dominant Forces in Protein Folding", Biochemistry, 29, pp. 7133-7155 (1990)				
24	C. Dinarello, "Role of Interleukin-1 in Infectious Disease				
	C. Dinarello et al., "Anticytokine Strategies in the Treat Syndrome", J. Am. Med. Assoc., 269, pp. 1829-1835 (ment of the Systemic Inflammat			
	R. Dolle et al., "Aspartyl α-((Diphenylphosphinyl)oxy)m 1β Converting Enzyme. Utility of the Diphenylphosphin Cysteine Proteases", J. Med. Chem., 38, pp. 220-222 (ethyl Ketones as Novel Inhibitor nic Acid Leaving Group for the Ir			
H	R. Dolle et al., "Aspartyl α((1-Phenyl-3-(trifluoromethyl) 1β Converting Enzyme Inhibitors. Significance of the FInhibitor Binding", J. Med. Chem., 37, pp. 3863-3865 (1	P ₁ and P ₃ Amido Nitrogens for E			
J.J.	R. Dolle et al., "P $_1$ Aspartate-Based Peptide α -((2,6-Did Time-Dependent Inhibitors of Interleukin-1 β -Converting (1994)	chlorobenzoyl)oxy)methyl Keton g Enzyme" <u>J. Med. Chem.,</u> 37, p	es as Potent p. 563-564		
SHX	S. Ealick et al., "Application of Crystallographic and Mo Nucleoside Phosphorylase Inhibitors", Proc. Natl. Acad				
L	P. Edwards et al., "Design, Synthesis, and Kinetic Eval the Peptidyl α-Ketobenzoxazoles, and the X-ray Crysta Porcine Pancreatic Elastase and Ac-Ala-Pro-Val-2-Ber 1863 (1992)	al Structure of the Covalent Com	nplex between		
	H. Eklund et al "Three-dimensional Structure of Horse Resolution", J. Mol. Biol., 102, pp. 27-59 (1976)	e Liver Alcohol Dehydrogenase	at 2.4 Å		
EXAMINER	David Lutton	DATE CONSIDERED	6/13/03		

FORM PTO-1	U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY. DOCKET NO. VPI/94-04 CIP2 DIV5	SERIAL NO. 09/886,773		
(PE)	INFORMATION DISCLOSURE STATEMENT BY APPLICANT	APPLICANT Guy W. Bemis, et al.			
JAN 1 5 2002		FILING DATE June 21, 2001	GROUP 1653		
PER STRAINERS	OTHER DOCUMENTS (Including Author, Title,	Date, Pertinent Pages, Etc.)			
EXAMINER INITIAL			nije og gjer		
DL	P.R. Elford, et al., "Reduction of Inflammation and Pyre 015, an Inhibitor of the Interleukin-1β Converting Enzyr 601-606 (1995)				
) JY	J. Erickson et al., "Design, Activity, and 2.8 Å Crystal S to HIV-1 Protease", Science, 249, pp. 527-533 (1990)		oitor Complexed		
DL	TP.D. Fan et al., "Stimulation of Angiogenesis by Sul Inhibition by NK ₁ or Interleukin-1 Receptor Antagonists				
DL.	I. Fauszt et al., "Inhibition of Interleukin-1β Converting Am. Peptide Symp., June 20-25, 1993; Hodges, R.S. a (1994)				
	D.S. Fletcher, et al., "A Synthetic Inhibitor of Interleukir Induced Interleukin-1 <u>B</u> Production In Vitro and In Vivo" (1995)				
JL.	V. Gagliardini et al., "Prevention of Vertebrate Neuronal Death by the crmA Gene", <u>Science</u> , 263, pp. 826-828 (1994)				
	T. Geiger et al., "Neutralization of Interleukin-1β Activit Collagen-induced Arthritis in DBA/1 Mice and Prevents Exp. Rheumatol, 11, pp. 515-522 (1993)				
DY DY	A. Giannis & T. Kolter, "Peptidomimetics for Receptor Perspectives", Agnew. Chem. Int. Ed. Engl. 32, pp. 124		nt, and Medical		
YY ,	P. Goodford, "A Computational Procedure for Determine Biologically Important Macromolecules", J. Med. Chem		ding Sites on		
ZZ	D. Goodsell & A. Olson, "Automated Docking of Substi Proteins: Structure, Function, and Genetics, 8, pp. 195		nnealing",		
	OTHER DOCUMENTS (Including Author, Title, I	Date, Pertinent Pages, Etc.)			
EXAMINER INITIAL		* * * * *	a sugar series seguine		
De la companya della companya della companya de la companya della	T. Graybill et al., "The Preparation and Evaluation of Plantibitors of ICE", Am. Chem. Soc. Abs. (206th Natl. M		Reversible		
WY.	T. Graybill, et al., "Preparation and Evaluation of Peption of Interleukin-1β Converting Enzyme (ICE)", Int. J. Pep				
WK V	W. Griffin et al., "Brain Interleukin 1 and S-100 Immuno Alzheimer Disease", <u>Proc. Natl. Acad. Sci USA</u> , 86, pp		Syndrome and		
	C. Hammerberg et al., "interieukin- i Receptor Antagor Invest., 90, pp. 571-583 (1992)	nist in Normal and Psoriatic Epic	lermis", <u>J. Clin.</u>		
EXAMINER	David Latter	DATE CONSIDERED	6/13/03		

FORM PTO-1	U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY. DOCKET NO. VPI/94-04 CIP2 DIV5	SERIAL NO. 09/886,773		
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PEN & IRADE	OTHER DOCUMENTS (Including Author, Title, I	Date, Pertinent Pages, Etc.)			
EXAMINER INITIAL	经常数性处理的发表 。		COLORS		
DL.	S. Hanessian et al., "Design and Synthesis of a Prototy Receptor" <u>Bioorg. Med. Chem. Lett.</u> , 4, 1397-1400 (19		inin NK-2		
	E. Harris, "Rheumatoid Arthritis: Pathophysiology and pp. 1277-1289 (1990)		g. J. Med., 322,		
OL .	W. Hendrickson et al., "Selenomethionyl Proteins Prod Diffraction (MAD): A Vehicle for Direct Determination of 1665-1672 (1990)				
DL	R. Hirschmann et al., "The First Design and Synthesis of a Steroidal Peptidomimetic. The Potential Value of Peptidomimetics in Elucidating the Bioactive Conformation of Peptide Ligands", <u>J. Am. Chem.</u> Soc., 114, pp. 9699-9701 (1992)				
9£	R. Hirschmann et al., "Nonpeptidal Peptidomimetics with a β-D-Glucose Scaffolding. A Partial Somatostatin Agonist Bearing a Close Structural Relationship to a Potent, Selective Substance P Antagonist", J. Am. Chem. Soc., 114, pp. 9217-9218 (1992)				
	A. Holmgren et al., "Three-dimensional Structure of Escherichia coli Thioredoxin-S ₂ to 2.8Å Resolution", Proc. Natl. Acad. Sci. USA, 72, pp. 2305-2309 (1975)				
YX	A. Hopfinger, "Computer-Assisted Drug Design", J. Me	d. Chem., 28, pp. 1133-1139 (1	985)		
L	A. Hopfinger & B. Burke, "Molecular Shape Analysis: A Molecular Similarity", Concepts and Applications of Molegiora eds., 1990)				
DL _	A. Howard et al., "High-Level Production and Characte Converting Enzyme in Baculovirus and E.coli Expressi p. 146 (1993)				
W.L	A. Howard et al., "Human Interleukin-1β Converting En Activation", <u>J. Cell. Biochem. Suppl.</u> , 17B, p. 113 (1993)		Proenzyme		
	A. Howard et al., "IL-1-Converting Enzyme Requires A Precursor at Two Distinct Sites and Does Not Cleave 3 (1991)				
D DY	I. Kamphuis et al., "Thiol Proteases: Comparative Stud Papain and Actinidin, and on Amino Acid Sequence In Bromelain", <u>J. Mol. Biol.</u> , 182, pp. 317-329 (1985)				
	J. Knowles, "Tinkering with Enzymes: What are We Le	arning?", <u>Science</u> , 236, pp. 125	2-1258 (1987)		
	M. Kostura et al., "Identification of a Monocyte Specific Natl. Acad. Sci. USA, 86, pp. 5227-5231 (1989)	Pre-Interleukin 1β Convertase	Activity", <u>Proc.</u>		
ŽŽŽ	K. Kuida et al., "Altered Cytokino Export and Apoptosis Enzyme, Science, 267, pp. 2000-2003 (1995)	in Mice Deficient in Interleukin-	1β Converting		
EXAMINER	David Lister	DATE CONSIDERED	6/13/03		

FORM PTO-1	449 U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY. DOCKET NO. VPI/94-04 CIP2 DIV5	SERIAL NO. 09/886,773		
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TRAIL & LEADE	OTHER DOCUMENTS (Including Author, Title,	Date, Pertinent Pages, Etc.)	CE SE		
EXAMINER ANITIAL					
W.	I. Kuntz et al., "A Geometric Approach to Macromolect pp. 269-288 (1982)	ule-Ligand Interactions", <u>J. Mol.</u>	Biol., 16		
~ XX	E. Lattman, "Use of the Rotation and Translation Fund	ctions", Methods Enzymol., 115,	pp. 55-77 (1985)		
W.	P. Li et al., "Mice Deficient in IL-1β-Converting Enzyme Resistant to Endotoxic Shock", <u>Cell</u> , 80, pp. 401-411 (e are Defective in Production of	C, 27		
XX	C. Lipinski, "Bioisosterism in Drug Design", Annu. Rep	o. Med. Chem., 21, pp. 283-291	(1986)		
	G. Lonnemann et al., "Differences in the Synthesis and Kinetics of Release of Interleukin 1β and Tumor Necrosis Factor from Human Mononuclear Cells", <u>Eur. J. Immunol.</u> , 19, pp. 1531-1536 (1989)				
	A. MacKenzie et al., "An Inhibitor of the Interleukin-1β-Processing Enzyme Blocks IL-1 Release and Reduces Pyrexia and Acute Inflammation", Inflammation Research Association (7th Internat. Conf.), W42 (1994)				
A	T. Mandrup-Poulsen et al., "Involvement of Interleukin β-Cell Destruction in Insulin-dependent Diabetes Melli				
	C. March et al., "Cloning, Sequence and Expression of Two Distinct Human Interleukin-1 Complementary DNAs", Nature, 315, pp. 641-647 (1985)				
XX ,	G. Marshall, "Computer-Aided Drug Design", Annu. Rev. Pharmacol. Toxicol., 27, pp. 193-213 (1987)				
Je De	G. Marshall & I. Motoc, "Approaches to the Conformation of the Drug Bound to the Receptor", Molecular Graphics and Drug Design, pp. 115-156 (A. Burgen et al. eds., 1986)				
62 () ()	Y. Martin, "3D Database Searching in Drug Design", J. Med. Chem., 35, pp. 2145-2154 (1992)				
A 12	J. Marx, "Cell Death Studies Yield Cancer Clues", Science, 259, pp. 760-761 (1993)				
/ *	D. Mayer et al., "A Unique Geometry of the Active Site with Structure-Activity Studies", J. Comput. Aided Mol.		zyme Consistent ——————		
XX	R. Ménard et al., "Contribution of the Glutamine 19 Sic Oxyanion Hole of Papain", <u>Biochemistry</u> , 30, pp. 8924		bilization in the		
AL N	R. Ménard et al., "Importance of Hydrogen-Bonding In the Catalytic Mechanism of Papain", Biochemistry, 30,		ain of Asp 158 in		
	E. Meng et al., "Automated Docking with Grid-Based Epp. 505-524 (1992)	Energy Evaluation", <u>J. Comput. C</u>	Chem., 13,		
AL .	B. Miller et al., "Inhibition of Mature IL-1 β Production in Inflammation by WIN 67694, an Inhibitor of IL-1 β Conv. 1338 (1995)				
W.	D. Miller et al., "The IL-1β Converting Enzyme as a Th 133-148 (1993)	erapeutic Target", Ann. N.Y. Ac	ad. Sci., 696, pp.		
EXAMINER	Quil 1	DATE CONSIDERED	6/13/03		

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ATTY. DOCKET NO.

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EXAMINER INITIAL	The second secon				
X	S. Miller et al., "The Accessible Surface Area and Sta 836 (1987)	ability of Oligomeric Proteins",	Nature, 328 pp. 834-		
	A. Miranker & M. Karplus, "Functionality Maps of Bind Method", Proteins: Structure, Function, and Genetics		nultaneous Search		
YL	M. Miura et al., "Induction of Apoptosis in Fibroblasts Homolog of the C. elegans Cell Death Gene ced-3", 9		a Mammalian		
DI DI	A.M.M. Mjalli et al., "Activated Ketones as Potent Reversible Inhibitors of Interleukin-1β Converting Enzyme", Bioorg. Med. Chem. Lett., 4, pp. 1965-1968 (1994)				
D'Z	A.M.M Mjalli. et al., "Phenylalkyl Ketones as Potent Reversible Inhibitors of Interleukin-1β Converting Enzyme," Bioorg. Med. Chem. Lett., 3, pp. 2689-2692 (1993)				
DL.	S. Molineaux et al., "Interleukin 1β (IL-1β) Processing in Murine Macrophages Requires a Structurally Conserved Homologue of Human IL-1β Converting Enzyme", Proc. Natl. Acad. Sci. USA, 90, pp. 1809-1813 (1993)				
H .	B. Mosley et al., "Determination of the Minimum Poly Human Interleukins 1α and 1β", Proc. Natl. Acad. Sc	peptide Lengths of the Functic ie. USA, 84, pp. 4572-4576 (1	nally Active Sites of 987)		
J. D.L.	M.D. Mullican et al., "The Synthesis and Evaluation of Peptidyl Aspartyl Aldehydes as Inhibitors of ICE" Bioorg. Med. Chem. Lett., 4, 2359-2364 (1994)				
AL A	C.M. Nalin, "Apoptosis Research Enters the ICE Age	," <u>Structure,</u> 3, pp. 143-145 (1	995)		
	M. Navia & M. Murcko, "Use of Structural Information pp. 202-210 (1992)	n in Drug Design", <u>Curr. Opin.</u>	Struc. Biol., 2,		
)L	M. Nett et al., "Molecular Cloning of the Murine IL-1β Converting Enzyme cDNA", <u>J. Immunol.</u> , 149, pp. 3254-3259 (1992)				
	M. Nett-Fiordalisi et al., "Characterization and Activat Enzyme", J. Cell. Biochem. Suppl., 17B, p. 117 (1993)		β (IL-1β) Converting		
HL .	Y. Nishibata & A. Itai, "Automatic Creation of Drug Candidate Structures Based on Receptor Structure. Starting Point for Artificial Lead Generation", Tetrahedron, 47, pp. 8985-8990 (1991)				
XX	C. Noren et al., "A General Method for Site-Specific Incorporation of Unnatural Amino Acids into Proteins, Science, 244, pp. 182-188 (1989)				
H.	I. Noronha et al., "In situ Production of TNF-α, IL-1β a Kidney Int., 43, pp. 682-692 (1993)	and IL-2R in ANCA-positive G	omerulonephritis",		
a St	K. Ohlsson et al., "Interleukin-1 Receptor Antagonist 348, pp. 550-552 (1990)	Reduces Mortality from Endot	oxin Shock", Nature,		
XX	J. Oppenheim et al., "There is More than One Interley	ıkin 1" İmmunol Today 7 nr	45 56 (1986)		

U.S. DEPARTMENT OF COMMERCE

FORM PTO-1449

EXAMINER

EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not conformance and not considered. Include copy of this form with next communication to applicant.

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	OTHER DOCUMENTS (Including Author, Title,	Date, Pertinent Pages, Etc.)	CENT		
EXAMINER IŅITIAL					
W.	M. Pennington & N. Thornberry, "Synthesis of a Fluoro Substrate Based on Resonance Energy Transfer", Per	genic Interleukin-1β Converting ot. Res., 7, pp. 72-76 (1994)	Enzyme 0		
J. DK	R. Peters & R. McKinstry, "Three-Dimensional Modelin Design Arrived?" <u>Biotechnology</u> , 12, pp. 147-150 (1994)	ng and Drug Development: Has ' 4)	'Rational" Drug		
XX and	J. Plattner & D. Norbeck, "Obstacles to Drug Developn Technologies, pp. 92-126 (C. Clark & W. Moss eds., 1		ug Discovery		
	L. Polgár, "On the Mode of Activation of the Catalytical Biochem., 33, pp. 104-109 (1973)	ly Essential Sulfhydryl Group of	Papain", <u>Eur. J.</u>		
DL	C. Prasad et al., "P ₁ Aspartate-Based Peptide α-Arylad Time-Dependent Inhibitors of Interleukin-1β Converting Chem. Symp.), 66 (1994)				
	C. Ray et al., "Viral Inhibition of Inflammation: Cowpox Converting Enzyme", Cell, 69, pp. 597-604 (1992)	Virus Encodes an Inhibitor of th	e Interleukin-1β		
\mathcal{X}	L. Reiter, "Peptidic p-Nitroanilide Substrates of Interleu Res., 43, pp. 87-96 (1994)	ıkin-1β-Converting Enzyme", <u>Int.</u>	J. Pept. Protein		
	L. Revesz et al., "Synthesis of P1 Aspartate-Based Peptide Acyloxymethyl and Fluoromethyl Ketones as Inhibitors of Interleukin-1β-Converting Enzyme", <u>Tetrahedron Lett.</u> , 35, pp. 9693-9696 (1994)				
	C. Ring et al., "Structure-based Inhibitor Design by Using Protein Models for the Development of Antiparasitic Agents", Proc. Natl. Acad. Sci. USA, 90, pp. 3583-3587 (1993)				
	R.P. Robinson and K.M. Donahue, "Synthesis of a Pep Side Chain: An Inhibitor of Interleukin-1β Converting E				
\mathcal{A}	M.J. Salvatore et al., "L-741,494, A Fungal Metabolite Enzyme," J. Nat. Prods., 57, 755-760 (1994)	that is an Inhibitor of Interleukin	-1β Converting		
	J. Sandberg et al., "Treatment with an Interleukin-1 Re Allograft Survival", <u>Diabetes</u> , 42, pp. 1845-1851 (1993)		ngs Mouse Islet		
OL L	I. Schechter & A. Berger, "On the Size of the Active Sit Res. Commun., 27, pp. 157-162 (1967)	e in Proteases. I. Papain", <u>Bio</u>	chem. Biophys.		
	S. Schmidt et al., "Synthesis and Evaluation of Asparty α-Arylacyloxymethyl Ketones as Inhibitors of Interleuki (208th ACS Natl. Mtg.), MEDI 4, (1994)		Chem. Soc. Abs.		
	B. Shivers et al., "Molecular Cloning of Rat Interleukin-Regulation", J. Cell. Biochem. Suppl., 17B, p. 119 (199		tion and		
il il	 Singer et al., "Interleukin 1β is Localized in the Cytop from the Goigi Apparatus and Plasma Membranes of S pp. 389-407 (1988) 	lasmic Ground Substance but is			
EXAMINER	David Letter	DATE CONSIDERED	6/13/03		

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VPI/94-04 CIP2 DIV5	09/886,773
APPLICANT Guy W. Bemis, et al.	
FILING DATE	GROUP
June 21, 2001	1653

- STRAIN	OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)
EXAMINER ,kŅITIAL	
De la companya della companya della companya de la companya della	P. Sleath et al., "Substrate Specificity of the Protease that Processes Human Interleukin-1β, <u>J. Biol.</u> Chem., 265, pp. 14526-14528 (1990)
	A.F. Spatola, in "Chemistry and Biochemistry of Amino Acids, Peptides, and Proteins,", 7, ch. 5, pp. 267-281, Weinstein, B., ed., Marcel Dekker, Inc., New York (1983)
SH ,	R. Taylor & O. Kennard, "Hydrogen-Bond Geometry in Organic Crystals", Acc. Chem. Res., 17, pp. 320-326 (1984)
	C. Thornber, "Isosterism and Molecular Modification in Drug Design", <u>Chem. Soc. Rev.</u> , 8, pp. 563-580 (1979)
	N. Thornberry et al., "A Novel Heterodimeric Cysteine Protease is Required for Interleukin-1β Processing in Monocytes", Nature, 356, pp. 768-774 (1992)
	N. Thornbery et al., "Inactivation of Interleukin-1β Converting Enzyme by Peptide (Acyloxy) methyl Ketones", <u>Biochemistry</u> , 33, pp. 3934-3940 (1994)
	J. Travis, "Proteins and Organic Solvents Make an Eye-Opening Mix", Science, 262, p. 1374 (1993)
W.Z	J. Uhl et al., "Secretion of Human Monocyte Mature IL-1β: Optimization of Culture Conditions and Inhibition by ICE Inhibitors", Inflammation Res., 44, pp. S211-S212 (1995)
YL .	P. Warner, et al., "PYI Idone HLE Inhibitors: Variation of the 3 and 5 Substituents", Royal Soc. Chem. Abs. (7th RSC-SCI Med. Chem. Symp.), P23 (1993)
	S. Weiner et al. "A New Force Field for Molecular Mechanical Simulation of Nucleic Acids and Proteins", J. Am. Chem. Soc., 106, pp. 765-784 (1984)
	C. Wong & J. McCammon, "Dynamics and Design of Enzymes and Inhibitors", <u>J. Am. Chem. Soc.</u> , 108, pp. 3830-3832 (1986)
	P. Wooley et al., "The Effect of an Interleukin-1 Receptor Antagonist Protein on Type II Collagen-induced Arthritis and Antigen-induced Arthritis in Mice", Arthritis Rheum., 36, pp. 1305-1314 (1993)
XX	J. Yuan et al., "The C. elegans Cell Death Gene ced-3 Encodes a Protein Similar to Mammalian Interleukn-1β-Converting Enzyme", Cell, 75, pp. 641-652 (1993)

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DATE CONSIDERED

6/13/03